DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration 21 CFR Parts 201, 606, and 610

[Docket No. 2002N-0204]

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	Certifler R. GOSMA

Bar Code Label Requirement for Human Drug Products and Biological Products; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a final rule that appeared in the **Federal Register** of February 26, 2004 (69 FR 9120). The document included typographical and inadvertent errors. This document corrects those errors.

FOR FURTHER INFORMATION CONTACT: Philip L. Chao, Office of Policy and Planning (HF-23), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–0587.

SUPPLEMENTARY INFORMATION: In FR Doc. 04–4249, appearing on page 9120 in the **Federal Register** of Thursday, February 26, 2004, the following corrections are made:

- 1. On page 9151, in the third column, the first sentence of the first full paragraph, is corrected to read "We estimate that the rule provides net benefits to society of \$4.3 billion to \$4.5 billion annually, depending on whether a discount rate of 3 percent or 7 percent is used."
- 2. On page 9167, in the first column, the first sentence under the heading "P. Small Business Analysis and Discussion of Alternatives" is corrected to

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read "For the reasons cited in the following paragraphs, the agency certifies that the final rule will not have a significant economic impact on a substantial number of small entities."

Dated: 3.31.09

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. 04-????? Filed ??-??-04; 8:45 am]

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